

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
23 August 2001 (23.08.2001)

PCT

(10) International Publication Number
WO 01/60411 A1

(51) International Patent Classification⁷: A61K 47/48,
C07K 14/59

(74) Agents: TOYAMA, Tsutomu et al.; Yokoyama Building
6th floor, 4-10, Higashi Nihonbashi 3-chome, Chuo-ku,
Tokyo 103-0004 (JP).

(21) International Application Number: PCT/JP01/01167

(22) International Filing Date: 19 February 2001 (19.02.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
2000-41303 18 February 2000 (18.02.2000) JP

(71) Applicants (*for all designated States except US*):
KANAGAWA ACADEMY OF SCIENCE AND TECH-
NOLOGY [JP/JP]; 2-1, Sakado 3-chome, Takatsu-ku,
Kawasaki-shi, Kanagawa 213-0012 (JP). MITSUBISHI
CHEMICAL CORPORATION [JP/JP]; 5-2, Marunouchi
2-chome, Chiyoda-ku, Tokyo 100-0005 (JP).

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ,
DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
HU, ID, IL, IN, IS, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO,
NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR,
TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM,
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,
IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF,
CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published:

— with international search report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(72) Inventor; and

(75) Inventor/Applicant (*for US only*): KONDO, Noriaki
[JP/JP]; C/O MITSUBISHI KASEI INSTITUTE OF LIFE
SCIENCES, LIMITED, 11, Minamiooya, Machida-shi,
Tokyo 194-8511 (JP).

(54) Title: PHARMACEUTICAL COMPOSITION, REAGENT AND METHOD FOR INTRACEREBRAL DELIVERY OF PHARMACEUTICALLY ACTIVE INGREDIENT OR LABELING SUBSTANCE

(57) Abstract: There are provided a pharmaceutical composition which can easily transport a pharmaceutically active ingredient into a brain, a reagent can easily transport a labeling substance into a brain, and a method for delivering a pharmaceutically active ingredient or a labeling substance from blood into a brain. The pharmaceutically active ingredient and the labeling substance are bonded to a hibernation-specific protein, and used as a conjugate. By administering the conjugate together with a thyroid hormone substance or a substance enhancing secretion and production thereof, the pharmaceutically active ingredient or the labeling substance is delivered into a brain.

WO 01/60411 A1

DESCRIPTION

PHARMACEUTICAL COMPOSITION, REAGENT AND METHOD FOR
INTRACEREBRAL DELIVERY OF PHARMACEUTICALLY ACTIVE
INGREDIENT OR LABELING SUBSTANCE

Technical Field

The present invention relates to a pharmaceutical composition, a reagent and a method for intracerebral delivery of a pharmaceutically active ingredient or a labeling substance from blood, and so forth.

Background Art

The hibernation-specific protein was found as a protein of which amount changes in association with hibernation, which exists in blood of chipmunks and so forth (J. Biol. Chem., 267, 473-478 (1992)). The hibernation-specific protein includes four kinds of proteins (HP-20, HP-25, HP-27 and HP-55). HP-20, HP-25 and HP-27 have similar structures containing a collagen-like sequence (repetitions of Gly-X-Y) in about 40 residues of N-terminal amino acids, and HP-55 belongs to the α 1-antitrypsin-like serpin superfamily, which shows inhibitory activity against elastase (Japanese Patent Application Laid-open (Kokai) No. 9-157299). These proteins are specifically expressed in livers of chipmunks and so forth, and secreted into blood. In blood, HP-20, HP-25 and HP-27 form a hetero trimer (HP-20c, molecular weight: about 80 kDa) by the collagen-

like triple helix structure and intermolecular disulfide bonds between the half-cystine residues existing at the both termini of the triple helix structure, and HP-55 (molecular weight: about 60 kDa) is non-covalently bonded to the hetero trimer to form a complex.

In the case of chipmunks, it is known that the amount of the hibernation-specific protein circannually changes in association with hibernation, and the amount thereof in blood increases in the non-hibernating period and remarkably decreases during hibernation. To the contrary, in cerebrospinal fluid (namely, inside of a brain), it remarkably increases during hibernation period. Furthermore, the hibernation-specific protein in the cerebrospinal fluid is dissociated into HP-20c and HP-55. In the non-hibernating time with little hibernation-specific protein in cerebrospinal fluid, the hibernation-specific protein level in cerebrospinal fluid is as low as 1/1000 to 1/2000 of the level in blood. Transportation of the hibernation-specific protein into a brain is induced by administration of thyroxine (10-250 $\mu\text{g/kg}$) to animals, and the hibernation-specific protein level in cerebrospinal fluid increases within 1 day after the administration and reach a 40 to 50 times higher level within several days after the administration. Furthermore, if the administration of thyroxine is stopped, it returns to the original level.

Disclosure of the Invention

When it is intended to administer a pharmaceutically active ingredient or a labeling substance into a brain, substances that can be administered by a simple method such as administration into blood are limited, because the substance transportation from blood into a brain is strictly restricted. If there is provided a method enabling administration of a pharmaceutically active ingredient or a labeling substance into a brain by a simple method, it will become easy to transport the pharmaceutically active ingredient or the labeling substance into a brain at an organism level, and it will be extremely useful for diagnosis of brain diseases, tests for effects of drugs on the cerebral nerve system, development of novel drugs effective for the cerebral nerve system, elucidation of cerebral functions and so forth.

An object of the present invention is to provide a pharmaceutical composition which can easily transport a pharmaceutically active ingredient into a brain, a reagent which can easily transport a labeling substance into a brain, and a method for delivering a pharmaceutically active ingredient or a labeling substance from blood into a brain.

The inventor of the present invention assiduously studied in order to achieve the aforementioned object. As a result, it has been found that, if chipmunk blood plasma containing a hibernation-specific protein is intravenously administered to a nonhibernater not

producing the hibernation-specific protein to which thyroxine is administered, the hibernation-specific protein level in the cerebrospinal fluid of the animal was increased, and that fragments of the proteins constituting the hibernation-specific protein complex are sufficient for the formation of the hibernation-specific protein complex. Thus, the present invention has been accomplished.

The present invention provides a pharmaceutical composition comprising a conjugate that comprises one or more pharmaceutically active ingredients bound to a hibernation-specific protein (hereinafter also referred to as "pharmaceutical composition of the present invention"), and a reagent comprising a conjugate that comprises one or more pharmaceutically active ingredients bound to a hibernation-specific protein (hereinafter also referred to as "reagent of the present invention").

Preferably, the pharmaceutical composition of the present invention and the reagent of the present invention further comprise a thyroid hormone substance or a substance enhancing secretion and production thereof.

In the pharmaceutical composition of the present invention or the reagent of the present invention, the hibernation-specific protein may be each of the hibernation-specific proteins or a fragment thereof, or a complex constituted by them, for example, a complex between HP-20c and HP-55, a complex between a fragment

of HP-20c and HP-55, a complex between HP-20c and a fragment of HP-55, a complex between a fragment of HP-20c and a fragment of HP-55 and so forth.

In the pharmaceutical composition of the present invention, the pharmaceutically active ingredient or the labeling substance may be bonded to either one or both of HP-20c or a fragment thereof and HP-55 or a fragment thereof.

The present invention also provides an agent for inducing delivery of an intracerebrally transferable protein or a conjugate that comprises one or more pharmaceutically active ingredients or one or more labeling substances bound to an intracerebrally transferable protein, into a brain, which comprises a thyroid hormone substance or a substance enhancing secretion and production thereof as an active ingredient (hereinafter also referred to as "delivery-inducing agent of the present invention").

The present invention further provides a method for delivering a pharmaceutically active ingredient or a labeling substance into a brain, which comprises binding the pharmaceutically active ingredient or the labeling substance to a hibernation-specific protein to obtain a conjugate, and administering the conjugate together with a thyroid hormone substance or a substance enhancing secretion and production thereof (hereinafter also referred to as "delivery method of the present invention").

Brief Explanation of the Drawings

Fig. 1 shows the results of Western blot using rabbit antibodies directed to each of HP-22, HP-25, HP-27 and HP-55, which represent HP in blood and cerebrospinal fluid after administration of chipmunk plasma to thyroxine-treated rats.

Fig. 2 shows the results of immunohistochemical analysis of HP taken up into cerebral choroid epithelium cells of thyroxine-treated chipmunks.

Fig. 3 shows structures of HP-20c and HP-55. In the figure, the lines connecting two of C indicate that cystine is formed between them, and CHO indicates addition of a sugar chain. The boxed portions indicate such portions that, even if they are deleted, the association ability of HP-20c and HP-55 is maintained. The structures of HP-27, HP-25, HP-20 and HP-55 shown in the figure correspond to the sequence position of amino acid residues 31-215 shown in SEQ ID NO: 1, the sequence position of amino acid residues 29-215 shown in SEQ ID NO: 2, the sequence position of amino acid residues 24-196 shown in SEQ ID NO: 3, and the sequence position of amino acid residues 25-413 shown in SEQ ID NO: 4, respectively.

Fig. 4 shows the partial structures of HP-20c and HP-55 having the association ability. In the figure, the lines connecting two of C indicate that cystine is formed between them. SH indicates a free sulfhydryl group, and CHO indicates addition of a sugar chain. The partial structures of HP-27, HP-25, HP-20 and HP-55

shown in the figure correspond to the sequence position of amino acid residues 76-205 shown in SEQ ID NO: 1, the sequence position of amino acid residues 76-205 shown in SEQ ID NO: 2, the sequence position of amino acid residues 61-186 shown in SEQ ID NO: 3, and the sequence position of amino acid residues 279-413 shown in SEQ ID NO: 4, respectively.

Best Mode for Carrying out the Invention

Hereafter, embodiments of the present invention will be explained in detail.

<1> Pharmaceutical composition of the present invention

The pharmaceutical composition of the present invention contains a conjugate that comprises one or more pharmaceutically active ingredients bound to a hibernation-specific protein (this will also be referred to simply as "drug conjugate" hereinafter).

In the present invention, the hibernation-specific protein (this is also abbreviated as "HP" hereinafter) means, for example, the four kinds of the hibernation-specific proteins specific for hibernation found in chipmunks, fragments thereof, or a complex constituted by those proteins or fragments thereof. The four kinds of protein are called HP-27, HP-25, HP-20 and HP-55, and their typical amino acid sequences are shown in SEQ ID NOS: 1-4, respectively. These proteins are not limited to those having the amino acid sequences shown in SEQ ID NOS: 1-4, and include allogeneic mutants thereof.

HP-27, HP-25 and HP-20 form a hetero trimer (HP-

20c, molecular weight: about 80 kDa) by the collagen-like triple helix structure and intermolecular disulfide bonds between the half-cystine residues existing at the both termini of the triple helix structure, and HP-55 (molecular weight: about 60 kDa) is non-covalently bonded to the hetero trimer to form a complex. The aforementioned fragments are not particularly limited, so long as they possess an ability to form such a complex and the complex has an ability to transfer from blood into a brain, and include those fragments of the aforementioned four kinds of the proteins, obtained by partial digestion of the proteins. When HP has any activity against an animal organism to which HP is administered, to affect the animal organism, it is effective to delete or inactivate an active center of HP. The ability to form a complex and the ability to transfer from blood into a brain can be evaluated by such methods as described in the Examples mentioned hereinafter.

The HP complex may be a complex between HP-20c and HP-55, or a complex in which either one or both of HP-20c and HP-55 are fragments thereof. A fragment of HP-20c means a complex in which at least one of the proteins constituting HP-20c is digested.

The pharmaceutically active ingredient is a substance that exerts a pharmaceutical action on brain cells, and is not particularly limited so long as it can transfer from blood into a brain as the conjugate that comprises one or more pharmaceutically active

ingredients bound to the HP complex, according to the present invention. The pharmaceutically active ingredient may be labeled by bonding the substance for labeling as mentioned later thereto. Examples of the action of the pharmaceutically active ingredient include treatment of diseases in a brain, and specific examples of the pharmaceutically active ingredient include antitumor (anticancer) agents, antibiotics, agents for treating autoimmune deficiency syndrome (AIDS), agents for treating neuropathic diseases such as Parkinson's disease, neuro degenerative disease, multiple sclerosis, Alzheimer's disease and depression, anodynes, migraine-treating agents, and epilepsy-treating agents.

Also, examples of the pharmaceutically active ingredient having such an action, used in the present invention, include macromolecular compounds, hydrophilic compounds, and compounds having a number of polar functional groups, which by themselves can not reach a brain from blood. Specifically, examples of the antitumor agents includes methotrexate, adriamycin, cisplatin, cyclophosphamide, etoposide, and carboplatin. Examples of the antibiotics include amphotericin B, and gentamycin. Examples of the agents for treating neuropathic diseases include neurotrophic factors such as NGF (nerve growth factor), BDNF (brain-derived neurotrophic factor), CNTF (ciliary neurotrophic factor), GDNF (glial cell line-derived neurotrophic factor), NT-3 (neurotrophin 3), NT-4 (neurotrophin 4), NT-5 (neurotrophin 5), bFGF (basic fibroblast growth factor),

aFGF (acidic fibroblast growth factor), and EGF (epithelial growth factor); nerve system-acting cytokines such as interferon α , interferon β , interferon γ , IL-1 (interleukin 1), IL-2 (interleukin 2), IL-3 (interleukin 3), IL-4 (interleukin 4), IL-5 (interleukin 5), IL-6 (interleukin 6), TNF (tumor necrosis factor), GM-CSF (granulocyte macrophage colony stimulating factor), G-CSF (granulocyte colony stimulating factor), M-CSF (macrophage colony stimulating factor), and PDGF (platelet-derived growth factor); antagonists and agonists of hormones and neurotransmitters, such as somatostatin, oxytocin, vasopressin, guaranine, VIP (vasoactive intestinal polypeptide), adrenocorticotrophic hormone, CCK (cholecystokinin), substance P, bombesin, motilin, glicentin, glucagon, glucagon-like peptide, dopamine decarboxylase, tricosanthin, and adenosine; and neurotransmitters such as dopamine, GABA (γ -aminobutyric acid), acetylcholine, serotonin, noradrenalin, and various neuropeptides. Examples of the AIDS-treating agents include nucleic acid derivatives having anti-HIV activity, such as AZT (azidethymidine), dideoxyinosine and dideoxycytosine.

Derivatives of any of the above-mentioned pharmaceutically active ingredients are also used as the pharmaceutically active ingredient in the present invention.

A conjugate that comprises one or more pharmaceutically active ingredients bound to HP can be formed by binding HP and the pharmaceutically active

ingredient(s) by a method generally known as a method for binding a protein and one or more pharmaceutically active ingredients. Binding mode and binding sites of HP to the pharmaceutically active ingredient(s) are not particularly limited so long as the formed conjugate can transfer from blood into a brain. The ability to transfer from blood into a brain can be evaluated by such a method as described in the Examples mentioned hereinafter. Specifically, binding of HP and the pharmaceutically active ingredient may be performed by introducing an appropriate functional group at a site not affecting the activity of the pharmaceutically active ingredient, if necessary, and using an appropriate crosslinker and the like. Examples of the crosslinker include ones forming a peptide bond such as EDAC (1-ethyl-3-(3-dimethylaminopropyl)carbodiimide), ones forming a disulfide bond such as SPDP (N-succinimidyl-3-(2-pyridylthio)propionate) and cystamine, but are not limited thereto. Also, protein crosslinkers having additional function such as spacer function may be used and examples thereof include glutaraldehyde, DIPS (4,4'-diisothiocyanostilbene-2,2'-disulfonic acid), EMCS (N-(E-maleimidocaproyloxy)succinimide), GMBS (N-(gamma-maleimidobutyloxy)succinimide), DSP (dithiobis(succinimidylpropionate) and DTBPA (4,4'-dithiobis-phenylazide). Order of the formation of HP complex and the binding of the pharmaceutically active ingredient is not also limited. For example, it is

possible to (1) bind a pharmaceutically active ingredient to a complex between HP-20c and HP-55, (2) bind a pharmaceutically active ingredient to HP-55, and then allow association of them with HP-20c, (3) bind a pharmaceutically active ingredient to HP-55, and then allow association of them with HP-20c, or the like. Furthermore, the pharmaceutically active ingredient may be bound to either one or both of HP-20c or a fragment thereof and HP-55 or a fragment thereof. As to the site to which the pharmaceutically active ingredient binds, of HP or a fragment thereof, because HP-20c and HP-55 forms a complex as described below, the site is preferable a site which does not inhibit binding of HP-20c or a fragment thereof and HP-55 or a fragment thereof and which is not exposed on a surface in the steric structure of the protein as much as possible. The site is preferably close to the binding site of HP-20c or a fragment thereof and HP-55 or a fragment thereof. Specifically, in the case of HP-20c, it is preferably C-terminal side from 71Cys in SEQ ID NO: 3, and 111Cys is particularly preferable because it is a free sulfhydryl group. In the case of HP-55, it is preferably C-terminal side from, for example, 278Met in SEQ ID NO: 4, more preferably from 379Gln to 413Gln. With respect to the ratio of the pharmaceutically active ingredient to be bound and HP or a fragment thereof, the pharmaceutically active ingredient may be bound to all of active groups of HP or a fragment thereof using a protein crosslinker or the like, or may be bound part of

the active groups. If it is desired to bind the pharmaceutically active ingredient to a certain site-specific active group, it can be performed, for example, by masking the same kind of active groups at the other sites. Two or more kinds of the pharmaceutically active ingredients may be bound to one HP or a fragment thereof. The pharmaceutical composition prepared by binding as mentioned above may be evaluated for its ability to transfer into a brain according to such a method as described in the Examples mentioned hereinafter, or may be evaluated for activity of the pharmaceutically active ingredient, whether or not the conjugate itself has an active form or the like according to known methods, to screen one which is the best as the pharmaceutical composition of the present invention.

The conjugate can be contained in the pharmaceutical composition of the present invention also includes one in which both of one or more pharmaceutically active ingredients and one or more substances for labeling as mentioned later are bound to one HP or a fragment thereof, one in which one or more pharmaceutically active ingredients are bound to one of proteins constituting the HP complex or fragments thereof and one or more substances for labeling bound to another of the proteins or fragments thereof, and one which is a combination thereof.

The pharmaceutical composition of the present invention preferably further contains a thyroid hormone substance or a substance enhancing secretion and

production thereof. In this specification, the thyroid hormone substance means a thyroid hormone, a substance exhibiting thyroid hormone-like action, or a structural analogue of a thyroid hormone.

The thyroid hormone is preferably thyroxine or triiodothyronine.

The substance exhibiting thyroid hormone-like action and the structural analogue of the thyroid hormone mean a substance exhibiting action equivalent to a thyroid hormone used in the present invention, and they are not particularly limited so long as they exert action of transferring a conjugate that comprises one or more pharmaceutically active ingredients bound to HP, from blood to a brain.

Examples of the substance enhancing secretion and production of the thyroid hormone substance include a substance enhancing production and secretion of a thyroid hormone in thyroid, and specific examples thereof include thyroid-stimulating hormone (TSH), thyrotropin-releasing hormone (TRH) and so forth.

The pharmaceutical composition of the present invention may contain a pharmaceutically acceptable carrier in addition to the drug conjugate and the thyroid hormone substance.

The administration method and the dose of the pharmaceutical composition of the present invention may be similar to the administration method and the dose of the drug conjugate as will be explained hereinafter concerning the delivery method of the present invention.

Dosage form and content of the drug conjugate of the pharmaceutical composition of the present invention can be suitably decided by those skilled in the art according to the administration method, the dose and so forth. If the pharmaceutical composition of the present invention contains the drug conjugate and the thyroid hormone substance or a substance enhancing secretion and production thereof, they do not necessarily need to be mixed and the pharmaceutical composition of the present invention include those wherein they are contained individually and combined.

<2> Reagent of the present invention

The reagent of the present invention contains a conjugate that comprises one or more labeling substances bound to a hibernation-specific protein (this will be also referred to simply as "labeled conjugate" hereinafter), and is used as an imaging agent for encephalography, for diagnosis of cerebral diseases, screening for elucidation of brain functions and so forth.

The hibernation-specific protein and the hibernation-specific protein complex may be the same as those described above for the pharmaceutical composition of the present invention.

The labeling substance is not particularly limited so long as it is a substance labeled with a substance usually used for labeling. Examples of the substance used for labeling include, for example, fluorescent substances, radioisotopes or the like, coloring matters,

biotin and so forth. Antibodies and so forth can also be used.

As for the conjugate that comprises one or more labeling substances bound to the hibernation-specific protein, the production method, the order of binding and the structure of the labeled conjugate may be similar to those described above for the drug conjugate of the pharmaceutical composition of the present invention.

The reagent of the present invention preferably further contains a thyroid hormone substance or a substance enhancing secretion and production thereof. The thyroid hormone or the substance enhancing secretion and production thereof similar to one described above for the pharmaceutical composition of the present invention can be used.

The reagent of the present invention may contain a solvent usually used for injections, physiologically and pharmaceutically acceptable carrier and so forth, in addition to the labeled conjugate and the thyroid hormone substance or the substance enhancing secretion and production thereof.

The administration method and the dose of the reagent of the present invention can be suitably selected according to purposes of screening and so forth.

The dosage form, the content of the labeled conjugate and so forth of the reagent of the present invention can be suitably selected by those skilled in the art according to the administration method, the dose, the purpose of screening and so forth. When the reagent

of the present invention contains the labeled conjugate and the thyroid hormone substance or the substance enhancing secretion and production thereof, they do not necessarily need to be mixed, and the reagent of the present invention include those wherein they are contained individually and combined.

<3> Delivery-inducing agent of the present invention

The delivery-inducing agent of the present invention contains a thyroid hormone substance or a substance enhancing secretion and production thereof as an active ingredient, and induces delivery of an intracerebrally transferable protein or a conjugate that comprises one or more pharmaceutically active ingredients or one or more labeling substances bound to an intracerebrally transferable protein, into a brain.

The thyroid hormone substance or the substance enhancing secretion and production thereof, the pharmaceutically active ingredient and the labeling substance may be similar to those described above for the pharmaceutical composition of the present invention and the reagent of the present invention.

The intracerebrally transferable protein means a protein that can transfer from blood into a brain upon administration of the thyroid hormone substance or the substance enhancing secretion and production thereof. The ability to transfer from blood into a brain can be evaluated by, for example, measuring amounts of intracerebrally transferable protein in the brain or

cerebrospinal fluid with or without administration of the thyroid hormone substance or the substance enhancing secretion and production thereof by using such a method as described in the Examples mentioned hereinafter.

Examples of the intracerebrally transferable protein include HP. The intracerebrally transferable protein may be a fragment so long as its transfer into a brain is not inhibited. Preparation of a conjugate that comprises one or more pharmaceutically active ingredients or one or more labeling substances bound to an intracerebrally transferable protein can be performed in the same manner as explained above for the preparation of the conjugate that comprises one or more pharmaceutically active ingredients or one or more labeling substances bound to HP or a fragment thereof as for the pharmaceutical composition of the present invention and the reagent of the present invention.

The delivery-inducing agent of the present invention may contain a physiologically and pharmaceutically acceptable carrier and so forth in addition to the thyroid hormone substance or the substance enhancing secretion and production thereof. The administration method and the dose of the delivery-inducing agent of the present invention may be similar to the administration method and the dose of the thyroid hormone substance or the substance enhancing secretion and production thereof explained hereinafter as for the delivery method of the present invention. Specifically, the dosage form, the content of the thyroid hormone

substance or the substance enhancing secretion and production thereof and so forth of the delivery-inducing agent of the present invention can be suitably selected by those skilled in the art according to the administration method, the dose, the purpose of screening and so forth.

<4> Delivery method of the present invention

The delivery method of the present invention is a method for delivering a pharmaceutically active ingredient or a labeling substance into a brain, which comprises binding the pharmaceutically active ingredient or the labeling substance to a hibernation-specific protein to obtain a conjugate, and administering the resultant drug or labeled conjugate together with a thyroid hormone substance or a substance enhancing secretion and production thereof.

The pharmaceutically active ingredient, the labeling substance, HP and the thyroid hormone substance or the substance enhancing secretion and production thereof may be similar to those explained for the pharmaceutical composition of the present invention and the reagent of the present invention.

The binding of the pharmaceutically active ingredient or the labeling substance to HP can be performed in the same manner as explained for the preparation of a conjugate that comprises one or more pharmaceutically active ingredients or one or more labeling substances bound to HP as for the

pharmaceutical composition of the present invention and the reagent of the present invention.

In the delivery method of the present invention, the expression of "administering the drug or labeled conjugate together with a thyroid hormone substance or a substance enhancing secretion and production thereof" means that the drug or labeled conjugate and the thyroid hormone substance or the substance enhancing secretion and production thereof are administered so that there are attained levels of the both sufficient for causing transfer of the conjugate into a brain, and the administration is not limited to simultaneous administrations of the drug or labeled conjugate and the thyroid hormone substance or the substance enhancing secretion and production thereof. Therefore, either of the administration of the drug or labeled conjugate and the administration of the thyroid hormone substance or the substance enhancing secretion and production thereof may precede the other, or the both may be performed simultaneously. Preferably, the thyroid hormone substance or the substance enhancing secretion and production thereof is administered precedently.

Although the administration method of the drug or labeled conjugate is not particularly limited, it is preferable to administer the conjugate into blood. The administration into blood can be attained by, for example, intravenous injection. Dose can be suitably decided by those skilled in the art according to the body weight, age, condition of an object of

administration, the kind of the pharmaceutically active ingredient or the labeling substance, the purpose of screening and so forth.

The administration method of the thyroid hormone substance or the substance enhancing secretion and production thereof also is not particularly limited so long as such a level of the thyroid hormone substance or the substance enhancing secretion and production thereof that the drug or labeled conjugate can transfer from blood into a brain. For example, it can be intravenously or intraperitoneally administered singly or intravenously as a pharmaceutical composition together with the drug conjugate. The dose of the thyroid hormone substance or the substance enhancing secretion and production thereof can be suitably decided by those skilled in the art according to the kind of the thyroid hormone substance or the substance enhancing secretion and production thereof, the body weight, age, condition of an object of administration and so forth.

The present invention is based on the following findings.

After chipmunk plasma containing HP (HP-containing plasma) was administered to a rat, a nonhibernater, that did not produce HP, HP could be detected in the blood of the animal, but HP could not be detected in the cerebrospinal fluid. After thyroxine (250 $\mu\text{g}/\text{kg}$) was administered to a rat every day for 1 week, or one day after thyroxine (100 $\mu\text{g}/\text{kg}$) was administered to a rat, increase of HP in the cerebrospinal fluid could be

detected within 2-3 hours after the administration of the HP-containing plasma in the same amount as that before the thyroxine treatment.

Further, when choroid plexus epithelium cells of chipmunk in which intracerebral HP was increased by thyroxine treatment (10-250 $\mu\text{g/kg/day}$) were immunohistochemically stained by using an anti-HP-20c antibody, a strong positive reaction to the anti-HP-20c antibody was detected in the cytoplasm. This demonstrated that HP in blood was transported into the cerebrospinal fluid through choroid plexus, which is the blood-cerebrospinal fluid barrier.

The above observations means an unexpected result that HP, which is a heterogenous protein for a rat, was transported from blood into cerebrospinal fluid (into a brain) through the blood-cerebrospinal fluid barrier in the rat.

In order to determine a structure of HP effective for the transportation, structures of HP-20c and HP-55 having the ability to form a complex were investigated. As a result, HP-20c maintained the association ability with HP-55, even if about 40 of its N-terminus residues and about 10 of its C-terminus residues (boxed portions in Fig. 3) were partially deleted, and HP-55 could associate with HP-20c if about 150 residues from Met255 (this numbering corresponds to that used in the amino acid sequence shown in Fig. 3 (this residue corresponds to the amino acid number 279 in SEQ ID NO: 4)) to the C-terminus side (sequence shown in Fig. 3 except for the

boxed portion) were present.

From the above results, it was found that HP-20c and HP-55 could form a complex if they had the structures mentioned above. Therefore, since it is not considered that conservation of the whole structure of HP is essential for the transportation from blood into a brain, it is concluded that HP can be modified without affecting the transfer from blood into a brain.

Based on the above results, there was established a system for transporting a pharmaceutically active ingredient or a labeling substance from blood into a brain after administrations of a conjugate that comprises one or more pharmaceutically active ingredients or one or more labeling substances bound to HP, and the thyroid hormone substance into blood.

By using the method of the present invention utilizing HP and the thyroid hormone or the substance enhancing secretion and production thereof, intracerebral administration of an arbitrary pharmaceutically active ingredient or an arbitrary labeling substance can be artificially controlled. That is, the method of the present invention makes it possible to transport an arbitrary desired pharmaceutically active ingredient or an arbitrary desired labeling substance into a brain at an arbitrary particular time by administering the pharmaceutically active ingredient or the labeling substance bound to HP into blood and utilizing activation of the transportation ability from blood into a brain caused by

a thyroid hormone substance. Further, since HP forms a complex in blood and dissociates in a brain, it can also be realized that a pharmaceutically active ingredient and so forth delivered by this method should not exert its efficacy when it exists in blood, but can exert the efficacy only in a brain.

Example

Hereafter, the present invention will be further specifically explained with reference to the following example.

[Example 1]

(1) Transportation of HP in blood into rat cerebrospinal fluid

To a 12-week old rat, 0.4 ml of plasma collected from an adult chipmunk of a non-hibernating period (containing a complex between HP-20c and HP-55, referred to as "HP-containing plasma" hereinafter) was intravenously administered. Blood and cerebrospinal fluid were taken 2-3 hours after the administration, and HP in these samples was detected by Western blot using anti-HP antibodies (rabbit antibodies each directed to HP-20, HP-25, HP-27 and HP-55). The plasma in an amount of 0.13 μ l and the cerebrospinal fluid in an amount of 20 μ l were used for analysis. As a result, HP could be detected in the blood, but HP could not be detected in the cerebrospinal fluid (Fig. 1, a). After thyroxine was intraperitoneally administered to a rat (250 μ g/kg) every day for 1 week, the HP-containing plasma was

administered to the rat in the same manner as above, and HP in the blood and cerebrospinal fluid was detected. As a result, increase of HP could be detected in the cerebrospinal fluid within 2 to 3 hours after the administration. The results obtained after 2 hours are shown in Fig. 1, b. When the HP-containing plasma was administered once to a rat in the same manner as above 1 day after intraperitoneal administration of thyroxine (100 $\mu\text{g}/\text{kg}$), the same results were obtained.

(2) Mechanism of transportation

Thyroxine was intraperitoneally administered to an adult chipmunk at a non-hibernating period at a dose of 10 to 250 $\mu\text{g}/\text{kg}/\text{day}$ for six days. Intracerebral choroid plexus epithelium cells were isolated from this thyroxine-treated chipmunk. This sample was immunohistochemically stained by using an anti-HP-20c antibody. As a result, a strong positive reaction to the anti-HP-20c antibody was detected in the cytoplasm (Fig. 2). This demonstrated that HP in blood was transported into the cerebrospinal fluid through choroid plexus, which is the blood-cerebrospinal fluid barrier.

(3) Determination of modifiable site of HP

The portions for ability to form a complex were investigated in the structures of HP-20c and HP-55. The complex between HP-20c and HP-55 dissolved in physiological saline (phosphate buffered saline, PBS) was mixed with collagenase in a weight ratio of 2:1, and

incubated at 37°C for 3 hours. The digestion products were separated by gel permeation chromatography (GPC) to isolate the partially digested HP (C-HP). C-HP in an amount of 4 mg was mixed with 1 mg of lysyl endopeptidase, and incubated at 37°C for 5 hours for further partial digestion. The digestion products were separated by GPC to isolate the partially digested HP forming the complex. The obtained partially digested HP was further digested with lysyl endopeptidase in 50 mM Tris/hydrochloric acid buffer (pH 8.5) in the presence of 4 M urea, and the digestion products were separated by reverse phase HPLC. The peaks were each collected and attributed to peptides of HP-20c and HP-55 by comparison with peptide maps of HP-20c and HP-55 prepared similarly to determine the portions having the ability to form the complex.

As a result, HP-20c maintained the association ability with HP-55, even if about 40 of its N-terminus residues and about 10 of its C-terminus residues (boxed portion in Fig. 3) were partially deleted, and HP-55 could associate with HP-20c if about 150 residues from Met255 (this numbering corresponds to that used in the amino acid sequence shown in Fig. 3 (this residue corresponds to the amino acid number 279 in SEQ ID NO: 4)) to the C-terminus side (sequence shown in Fig. 3 except for the boxed portion) were present. From the above results, it was found that HP-20c and HP-55 could form a complex if they had the partial structures shown in Fig. 4.

From the above results, there was established a system for transporting a pharmaceutically active ingredient or a labeling substance from blood into a brain by administrations of a conjugate that comprises one or more pharmaceutically active ingredients or one or more labeling substances bound to HP, and a thyroid hormone substance or a substance enhancing secretion and production thereof into the blood.

Industrial Applicability

As explained above, the present invention provides a pharmaceutical composition which can easily transport a pharmaceutically active ingredient into a brain, and a reagent which can easily transport a labeling substance into a brain. These can be effective means for treatment, prevention and diagnosis of cerebral diseases and so forth. Further, according to the present invention, a pharmaceutically active ingredient or a labeling substance can be easily delivered into a brain by administration into blood. This further makes it easy to deliver various substances into a brain at an organism level, and provides novel means for investigations including elucidation of effects of drugs on the cerebral nerve system and elucidation of cerebral functions, development of novel drugs effective for the cerebral nerve system and so forth.

CLAIMS

1. A pharmaceutical composition comprising a conjugate that comprises one or more pharmaceutically active ingredients bound to a hibernation-specific protein.

2. The pharmaceutical composition according to Claim 1, which further comprises a thyroid hormone substance or a substance enhancing secretion and production thereof.

3. The pharmaceutical composition according to Claim 1 or 2, wherein the hibernation-specific protein is a complex between HP-20c and HP-55.

4. The pharmaceutical composition according to Claim 1 or 2, wherein the hibernation-specific protein is a complex between a fragment of HP-20c and HP-55, a complex between HP-20c and a fragment of HP-55, or a complex between a fragment of HP-20c and a fragment of HP-55.

5. The pharmaceutical composition according to Claim 3 or 4, wherein the pharmaceutically active ingredient is bonded to either one or both of HP-20c or a fragment thereof and HP-55 or a fragment thereof.

6. A reagent comprising a conjugate that comprises

one or more labeling substances bound to a hibernation-specific protein.

7. The reagent according to Claim 6, which further comprises a thyroid hormone substance or a substance enhancing secretion and production thereof.

8. The reagent according to Claim 6 or 7, wherein the hibernation-specific protein is a complex between HP-20c and HP-55.

9. The reagent according to Claim 6 or 7, wherein the hibernation-specific protein is a complex between a fragment of HP-20c and HP-55, a complex between HP-20c and a fragment of HP-55, or a complex between a fragment of HP-20c and a fragment of HP-55.

10. The reagent according to Claim 8 or 9, wherein the labeling substance is bonded to either one or both of HP-20c or a fragment thereof and HP-55 or a fragment thereof.

11. An agent for inducing delivery of an intracerebrally transferable protein or a conjugate that comprises one or more pharmaceutically active ingredients or one or more labeling substances bound to an intracerebrally transferable protein, into a brain, which comprises a thyroid hormone substance or a substance enhancing secretion and production thereof as

an active ingredient.

12. A method for delivering a pharmaceutically active ingredient or a labeling substance into a brain, which comprises binding the pharmaceutically active ingredient or the labeling substance to a hibernation-specific protein to obtain a conjugate, and administering the conjugate together with a thyroid hormone substance or a substance enhancing secretion and production thereof.

1/4

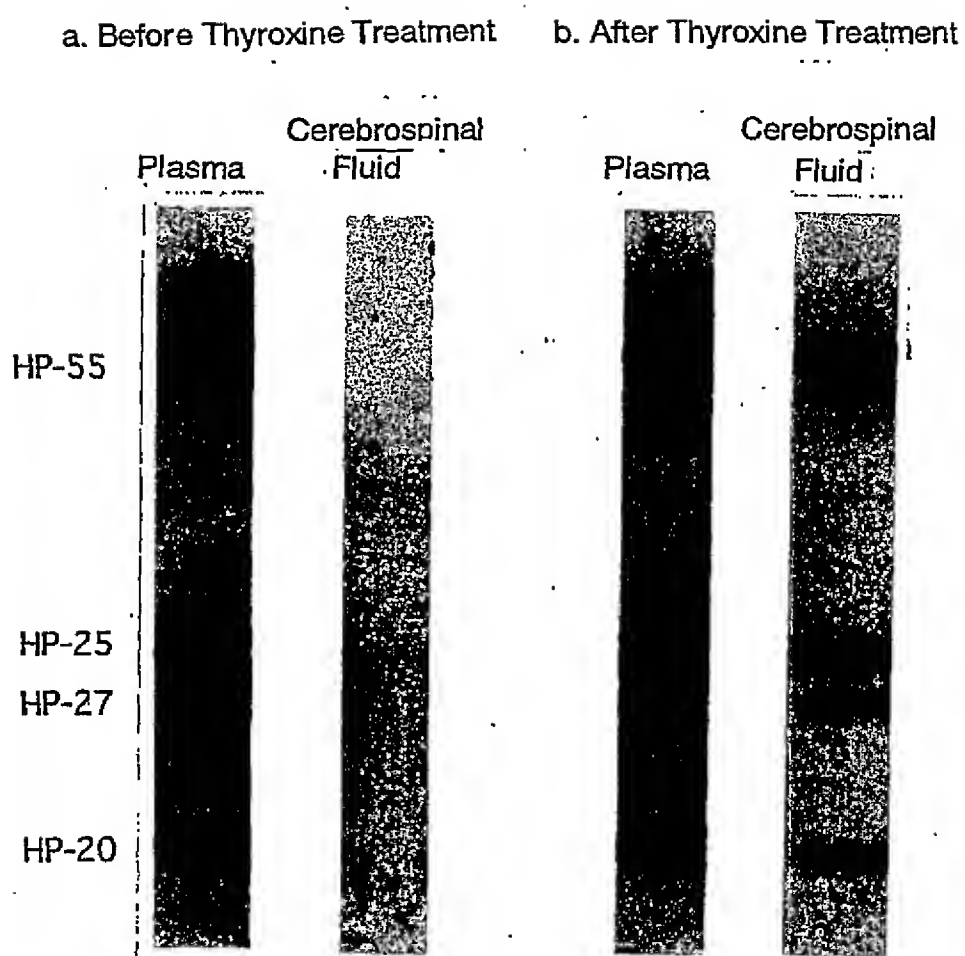
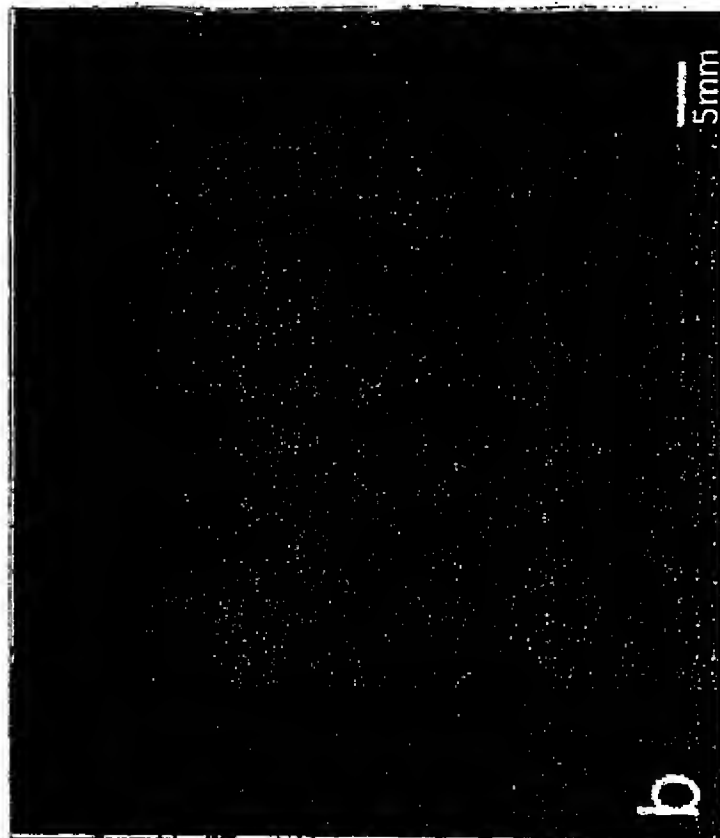


Fig. 1

Fluorescence Staining
Before Reaction with Anti-HP20c Antibody



Fluorescence Staining
After Reaction with Anti-HP20c Antibody



Fig. 2

3/4

Structure of HP-20c

HP-27 : ETQGNPES CNAPGPQGGP GMQGGPGTPG KPGPPGWNFG PGLPGPP GPP
 HP-25 : DSNNQGNSEP CGPPGPPGPP GIPGFPAGP ALGPPGPPGV PGIPGPQ GPP
 HP-20 : SGPPGPGVGP GVPVPGPRG PPGQGAAGR PGDPGPK GPS

GMTVNCHSKG TSAFAVKANE LPPAPSQPVI PKEALHDAQG HFDLATGVFT
 GDVEKCSSRP KSAFAVKLSE RPPEFFQPIV PKEALYNQEG HFNMATGEFS
 V KCPCRE RSAFTVKFSG RLPPPSEPVV FTEVLYNTQR DLKESTGVFN

SH CHO
 CPVPGLYQFG FHIEAVQRAV KVSIMRNGTQ VMEREABAQD GYEHISGTAI
 SH CHO
 CVLPGVYHFG FDIRLFQSSV KIRLMRDGIQ VREKRAQAND SYKRAMGSKI
CVEPGNYHFS EDVELYHCKV XIGLMENHIQ VMEKHQLSKN EYENASGAMI

LQLGMEDRVH LENKLSQTDL ERGTVOAVFS GFLIHEN
 MALGKGDKVH LESKLKGTES EKGITHVFF GYLLYKG
 MPLRQGDVW LEADVTEEP DQAKVVYFS GFLISS

Structure of HP-55

QDAQETEASK QDQHPASHR IAPHLAEFAL SLYRVLARQS NTFNIFFSRV
 SIASALAMLS LGTKGDTHTQ ILEGLDFNLT EMABADIHQG FQNLLOTLNR
 PNTQLQLTSG NVLFIHQNLK LLDKFLENIX SLYHSEAFPT NTFNMEEARQ
 QINSYVEKGT QGKIVELVKE LDSDTVLAIV NYIFFKKGWL KPFNVKNIRE
 EDFHVDEATT VRVPMYRVG MFPVHYCTRL ASLVLQMDYL GNATAIFLLP
 DKGRMOHLED TISTEILSKL LKDRQTSKYQ VYFPRVSISG TYDLKDVLS
 255
 LGITRVFSRV ADLSGVTEDA PLTVSKVLHK AVLDMDDEGT RAAGGTVLGA
 EAMLQAPIMK FDRPFLVVIY EHNTKSPLFV GKVVNPTQQ

Fig. 3

4/4

Partial Structure of HP-20c

HP-27 : GPP
 HP-25 : GPP
 HP-20 : GPS

GMTVNCHSKG TSAFAVKANE LPPAPSQPVI FKEALHDAQG HFDLATGVFT
 GDVEKCSRP KSAFAVKLSE RPPEPFQPIV FKEALYNQEG HFNMATGEFS
 V KCPCRE RSAFTVKFSG RLPPPSEPVV FTEVLYNTQR DLKESTGVFN
 SH CHO
 CPVPGLYQFG FHIEAVQRAV KVSLMRNGTQ VMEREARAQD GYEHISGTAI
 SH CHO
 CVLPGVYNFG FDIRLFQSSV KIRLMRDGIQ VREKEAQAND SYKHAMGSVI
CVEPGNYHFS FDVELYHCKV KIGLMKNHIQ VMEKHQLSKN EYENASGAMI
 LQLGMEDRVH LENKLSQTDL ERGTVQA
 MALGKGDKVW LESKLGTES EKGITHI
 MPLRQGDVW LEADVETEEP DQAKVV

Partial Structure of HP-55

MQHLED TISTEILSKL LKDRQTSKIYQ VYFPRVSISG TYDLKDVLS
 LGITRVFSRV ADLSGVTEDA PLTVSKVLHK AVLDMDDEGT EAAGGTVLGA
 EAMLQAPIMK FDRPFLVVYI EHNTKSPLFV GKVVNPTQQ

Fig. 4

1/5

Sequence Listing

<110> Kanagawa Academy of Science and Technology
Mitsubishi Chemical Corporation

<120> Pharmaceutical Composition, Reagent and Method for Intracerebral Delivery of Pharmaceutically Active Ingredient or Labeling Substance

<130> OP1157-PCT

<150> JP 2000-41303

<151> 2000-02-18

<160> 4

<210> 1

<211> 215

<212> PRT

<213> *Tamias asiaticus*

<400> 1

Met	Tyr	Glu	Ala	Gly	Lys	Arg	Ala	Ser	Phe	Met	Gly	Gly	Ala	Gly	Ile
1				5					10					15	
Trp	Ile	Leu	Ala	Leu	Ser	Val	Leu	Met	His	Val	Val	Cys	Ser	Glu	Thr
			20					25					30		
Gln	Gly	Asn	Pro	Glu	Ser	Cys	Asn	Val	Pro	Gly	Pro	Gln	Gly	Pro	Pro
		35					40					45			
Gly	Met	Arg	Gly	Pro	Pro	Gly	Thr	Pro	Gly	Lys	Pro	Gly	Pro	Pro	Gly
	50					55					60				
Trp	Asn	Gly	Phe	Pro	Gly	Leu	Pro	Gly	Pro	Pro	Gly	Pro	Pro	Gly	Met
	65				70				75					80	
Thr	Val	Asn	Cys	His	Ser	Lys	Gly	Thr	Ser	Ala	Phe	Ala	Val	Lys	Ala
			85					90					95		
Asn	Glu	Leu	Pro	Pro	Ala	Pro	Ser	Gln	Pro	Val	Ile	Phe	Lys	Glu	Ala
		100					105					110			
Leu	His	Asp	Ala	Gln	Gly	His	Phe	Asp	Leu	Ala	Thr	Gly	Val	Phe	Thr
		115				120						125			
Cys	Pro	Val	Pro	Gly	Leu	Tyr	Gln	Phe	Gly	Phe	His	Ile	Glu	Ala	Val

2/5

130	135	140
Gln Arg Ala Val Lys Val Ser Leu Met Arg Asn Gly Thr Gln Val Met		
145	150	155
Glu Arg Glu Ala Glu Ala Gln Asp Gly Tyr Glu His Ile Ser Gly Thr		160
	165	170
Ala Ile Leu Gln Leu Gly Met Glu Asp Arg Val Trp Leu Glu Asn Lys		175
	180	185
Leu Ser Gln Thr Asp Leu Glu Arg Gly Thr Val Gln Ala Val Phe Ser		190
	195	200
Gly Phe Leu Ile His Glu Asn		205
210	215	

<210> 2

<211> 215

<212> PRT

<213> *Tamias asiaticus*

<400> 2

Met Pro Ala Gln Arg Gly Gly Ala Leu Ser Met Gly Ala Ala Gly Phe		
1	5	10
Trp Ile Leu Val Leu Ser Ile Thr Ser Ala Leu Ala Asp Ser Asn Asn		15
	20	25
Gln Gly Asn Ser Glu Pro Cys Gly Pro Pro Gly Pro Pro Gly Pro Pro		30
	35	40
Gly Ile Pro Gly Phe Pro Gly Ala Pro Gly Ala Leu Gly Pro Pro Gly		45
	50	55
Pro Pro Gly Val Pro Gly Ile Pro Gly Pro Gln Gly Pro Pro Gly Asp		60
65	70	75
Val Glu Lys Cys Ser Ser Arg Pro Lys Ser Ala Phe Ala Val Lys Leu		80
	85	90
Ser Glu Arg Pro Pro Glu Pro Phe Gln Pro Ile Val Phe Lys Glu Ala		95
	100	105
Leu Tyr Asn Gln Glu Gly His Phe Asn Met Ala Thr Gly Glu Phe Ser		110
	115	120
Cys Val Leu Pro Gly Val Tyr Asn Phe Gly Phe Asp Ile Arg Leu Phe		125
	130	135
Gln Ser Ser Val Lys Ile Arg Leu Met Arg Asp Gly Ile Gln Val Arg		140
145	150	155
		160

3/5

Glu Lys Glu Ala Gln Ala Asn Asp Ser Tyr Lys His Ala Met Gly Ser
 165 170 175
 Val Ile Met Ala Leu Gly Lys Gly Asp Lys Val Trp Leu Glu Ser Lys
 180 185 190
 Leu Lys Gly Thr Glu Ser Glu Lys Gly Ile Thr His Ile Val Phe Phe
 195 200 205
 Gly Tyr Leu Leu Tyr Gly Lys
 210 215

<210> 3

<211> 196

<212> PRT

<213> *Tamias asiaticus*

<400> 3

Met Thr Asp Val Trp Arg Leu Ala Ile Phe Val Leu Met Val Asn Val
 1 5 10 15
 Leu Asn Asp Gln Val Ser Cys Ser Gly Pro Pro Gly Pro Val Gly Tyr
 20 25 30
 Pro Gly Val Pro Gly Val Pro Gly Pro Arg Gly Pro Pro Gly Gln Pro
 35 40 45
 Gly Ala Ala Gly Arg Pro Gly Asp Pro Gly Pro Lys Gly Pro Ser Val
 50 55 60
 Lys Cys Pro Cys Arg Glu Arg Ser Ala Phe Thr Val Lys Phe Ser Gly
 65 70 75 80
 Arg Leu Pro Pro Pro Ser Glu Pro Val Val Phe Thr Glu Val Leu Tyr
 85 90 95
 Asn Thr Gln Arg Asp Leu Lys Glu Ser Thr Gly Val Phe Asn Cys Val
 100 105 110
 Glu Pro Gly Asn Tyr His Phe Ser Phe Asp Val Glu Leu Tyr His Cys
 115 120 125
 Lys Val Lys Ile Gly Leu Met Lys Asn His Ile Gln Val Met Glu Lys
 130 135 140
 His Gln Leu Ser Lys Asn Glu Tyr Glu Asn Ala Ser Gly Ala Met Ile
 145 150 155 160
 Met Pro Leu Arg Gln Gly Asp Lys Val Trp Leu Glu Ala Asp Val Glu
 165 170 175
 Thr Glu Glu Pro Asp Gln Ala Lys Val Val Ile Tyr Phe Ser Gly Phe

4/5

180
Leu Ile Ser Ser
195

185

190

<210> 4
<211> 413
<212> PRT
<213> *Tamias asiaticus*

<400> 4

Met	Pro	Ser	Ser	Ile	Ser	Trp	Gly	Leu	Leu	Leu	Leu	Ala	Ala	Leu	Ser
1				5				10						15	
Cys	Leu	Gly	Pro	Gly	Ser	Leu	Ala	Gln	Asp	Ala	Gln	Glu	Thr	Glu	Ala
			20					25						30	
Ser	Lys	Gln	Asp	Gln	Glu	His	Pro	Ala	Ser	His	Arg	Ile	Ala	Pro	His
		35					40					45			
Leu	Ala	Glu	Phe	Ala	Leu	Ser	Leu	Tyr	Arg	Val	Leu	Ala	Arg	Gln	Ser
	50					55					60				
Asn	Thr	Thr	Asn	Ile	Phe	Phe	Ser	Pro	Val	Ser	Ile	Ala	Ser	Ala	Leu
65				70					75					80	
Ala	Met	Leu	Ser	Leu	Gly	Thr	Lys	Gly	Asp	Thr	His	Thr	Gln	Ile	Leu
				85					90					95	
Glu	Gly	Leu	Asp	Phe	Asn	Leu	Thr	Glu	Met	Ala	Glu	Ala	Asp	Ile	His
			100					105					110		
Gln	Gly	Phe	Gln	Asn	Leu	Leu	Gln	Thr	Leu	Asn	Arg	Pro	Asn	Thr	Gln
		115					120					125			
Leu	Gln	Leu	Thr	Ser	Gly	Asn	Val	Leu	Phe	Ile	His	Gln	Asn	Leu	Lys
	130					135					140				
Leu	Leu	Asp	Lys	Phe	Leu	Glu	Asn	Ile	Lys	Ser	Leu	Tyr	His	Ser	Gly
145				150					155					160	
Ala	Phe	Pro	Thr	Asn	Phe	Thr	Asn	Thr	Glu	Glu	Ala	Arg	Gln	Gln	Ile
				165					170				175		
Asn	Ser	Tyr	Val	Glu	Gln	Gly	Thr	Gln	Gly	Lys	Ile	Val	Glu	Leu	Val
		180						185					190		
Lys	Glu	Leu	Asp	Arg	Asp	Thr	Val	Leu	Ala	Leu	Val	Asn	Tyr	Ile	Phe
	195						200					205			
Phe	Lys	Gly	Lys	Trp	Leu	Lys	Pro	Phe	Asn	Val	Lys	Asn	Ile	Arg	Glu
	210					215						220			

5/5

Glu	Asp	Phe	His	Val	Asp	Glu	Ala	Thr	Thr	Val	Arg	Val	Pro	Met	Met
225					230					235					240
Tyr	Arg	Val	Gly	Met	Phe	Pro	Val	His	Tyr	Cys	Arg	Thr	Leu	Ala	Ser
			245						250					255	
Leu	Val	Leu	Gln	Met	Asp	Tyr	Leu	Gly	Asn	Ala	Thr	Ala	Ile	Phe	Leu
			260					265					270		
Leu	Pro	Asp	Lys	Gly	Lys	Met	Gln	His	Leu	Glu	Asp	Thr	Ile	Ser	Thr
		275					280					285			
Glu	Ile	Leu	Ser	Lys	Leu	Leu	Lys	Asp	Arg	Gln	Thr	Ser	Lys	Tyr	Gln
	290						295				300				
Val	Tyr	Phe	Pro	Arg	Val	Ser	Ile	Ser	Gly	Thr	Tyr	Asp	Leu	Lys	Asp
305					310					315					320
Val	Leu	Ser	Ser	Leu	Gly	Ile	Thr	Arg	Val	Phe	Ser	Arg	Val	Ala	Asp
				325					330					335	
Leu	Ser	Gly	Val	Thr	Glu	Asp	Ala	Pro	Leu	Thr	Val	Ser	Lys	Val	Leu
			340					345					350		
His	Lys	Ala	Val	Leu	Asp	Met	Asp	Glu	Glu	Gly	Thr	Glu	Ala	Ala	Gly
		355					360					365			
Gly	Thr	Val	Leu	Gly	Ala	Glu	Ala	Met	Leu	Gln	Ala	Pro	Ile	Met	Lys
	370						375				380				
Phe	Asp	Arg	Pro	Phe	Leu	Val	Val	Ile	Tyr	Glu	His	Asn	Thr	Lys	Ser
385					390					395					400
Pro	Leu	Phe	Val	Gly	Lys	Val	Val	Asn	Pro	Thr	Gln	Gln			
				405					410						

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61K47/48 C07K14/59

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A61K C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

BIOSIS, EPO-Internal, MEDLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	KONDO N ET AL: "IDENTIFICATION OF NOVEL BLOOD PROTEINS SPECIFIC FOR MAMMALIAN HIBERNATION" JOURNAL OF BIOLOGICAL CHEMISTRY, vol. 267, no. 1, 1992, pages 473-478, XP002165020 ISSN: 0021-9258 cited in the application the whole document --- -/-	1-12

☒ Further documents are listed in the continuation of box C.

☐ Patent family members are listed in annex.

* Special categories of cited documents:

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

Z document member of the same patent family

Date of the actual completion of the international search

9 April 2001

Date of mailing of the international search report

16.05.01

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Döpfer, K-P

INTERNATIONAL SEARCH REPORT

national application No.
PCT/JP 01/01167

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
see FURTHER INFORMATION sheet PCT/ISA/210
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this International application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.1

Although claim 12 is directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.

Continuation of Box I.1

Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy